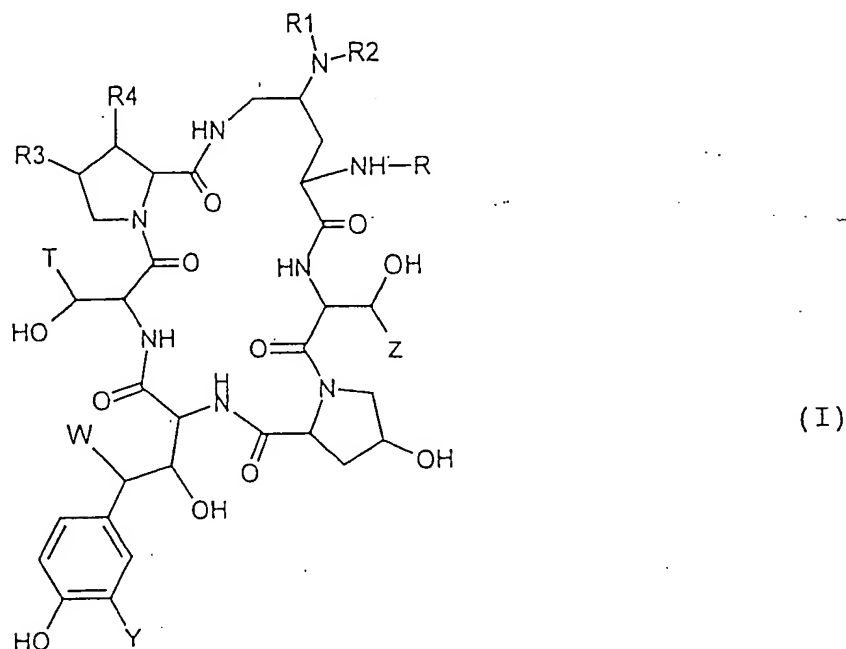


**Listing of Claims:**

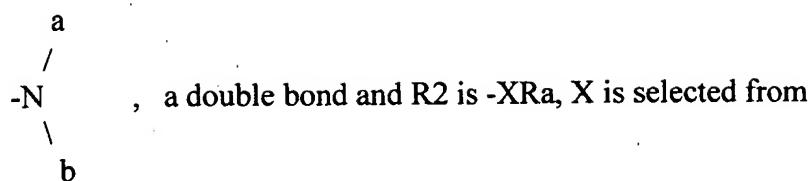
**Claim 1 (currently amended)** A compound selected from the group consisting of all possible isomeric forms and their mixtures of a compound of the formula



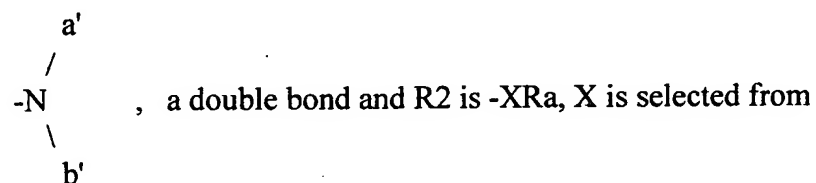
wherein either  $R_1$  and  $R_2$  are individually selected from the group consisting of hydrogen, hydroxyl, alkyl and cycloalkyl of up to 8 carbon atoms optionally interrupted by oxygen and optionally substituted by a member selected from the group consisting of



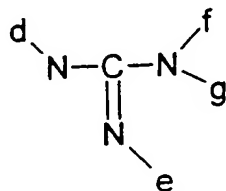
are individually hydrogen or alkyl of 1 to 8 carbon atoms ~~or a and b can optionally form with the nitrogen atom a heterocycle optionally containing at least one additional heteroatom~~, or  $R_1$  forms with the endocyclic carbon atom carrying



the group consisting of oxygen, -NH- or -N-alkyl of 1 to 8 carbon atoms and Ra is selected from the group consisting of hydrogen, alkyl or cycloalkyl of up to 8 carbon atoms substituted by at least one member of the group consisting of halogen, -OH, -CO<sub>2</sub>H, -CO<sub>2</sub>alk, and



a' and b' are hydrogen or alkyl of 1 to 8 carbon atoms ~~or a' and b' can form a heterocycle optionally containing at least one additional heteroatom or~~ and heterocycle containing at least one heteroatom or R<sub>2</sub> is



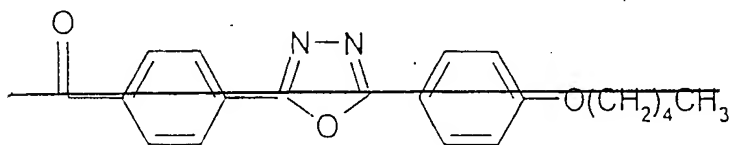
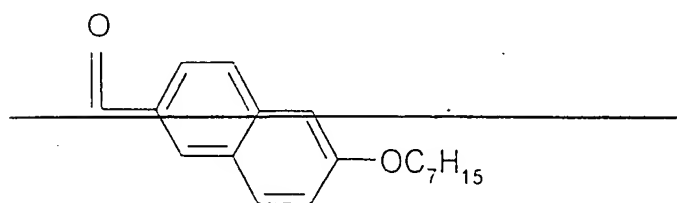
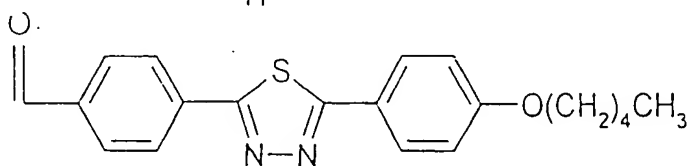
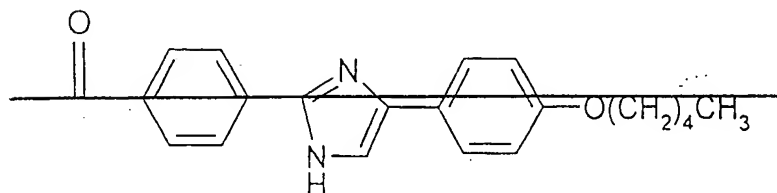
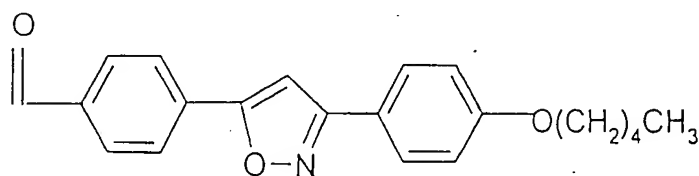
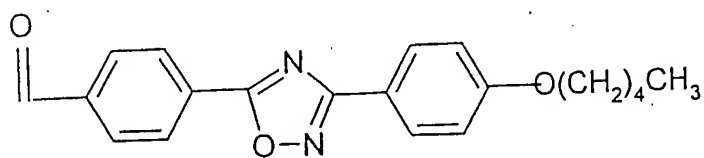
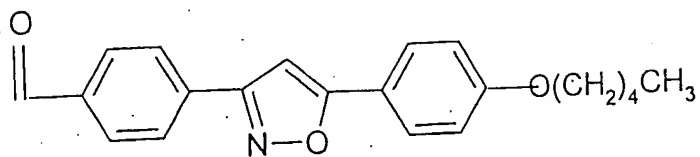
in which d, e, f and g are hydrogen or alkyl of 1 to 8 carbon atoms, f and g can also be acyl of up to 8 carbon atoms, ~~and e and f can also form a ring optionally containing at least one heteroatom,~~

R<sub>3</sub> is selected from the group consisting of hydrogen, methyl and hydroxyl,

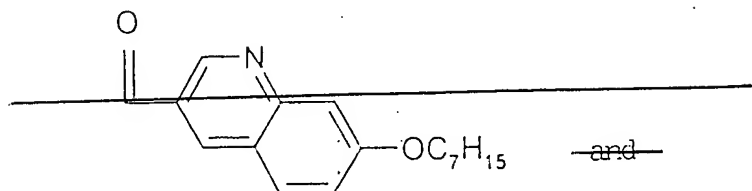
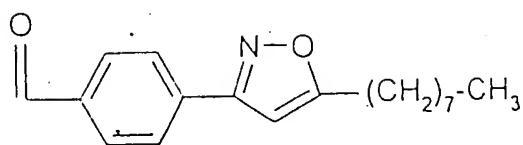
R<sub>4</sub> is hydrogen or hydroxyl,

R is selected from the group consisting of

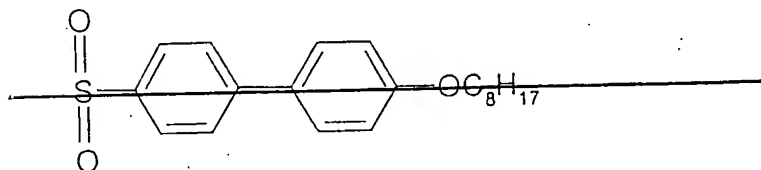
R is selected from the group consisting of



and



and



T is selected from the group consisting of hydrogen, methyl,  $-\text{CH}_2\text{CONH}_2$ ,  $-\text{CH}_2\text{CN}$ ,  $-(\text{CH}_2)_2\text{NH}_2$  and  $-(\text{CH}_2)\text{Nalk}^+\text{X}^-$ , X is halogen and alk is alkyl of up to 8 carbon atoms, Y is selected from the group consisting of hydrogen, hydroxyl, halogen and  $\text{OSO}_3\text{H}$  and a salt thereof, W is hydrogen or  $-\text{OH}$ , Z is hydrogen or methyl and a non-toxic, pharmaceutically acceptable acid addition salt thereof.

**Claim 2** (previously presented) A compound of claim 1 in which T is hydrogen.

**Claim 3** (previously presented) A compound of claim 1 in which W is hydrogen.

**Claim 4** (previously presented) A compound of claim 1 in which Z is methyl.

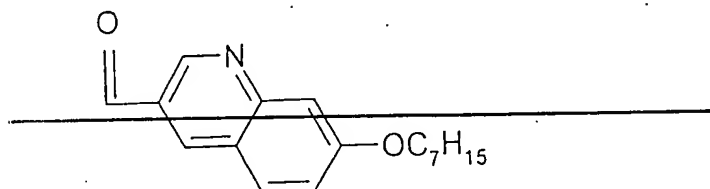
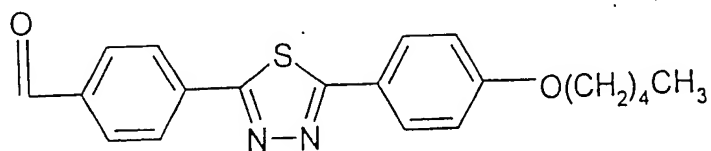
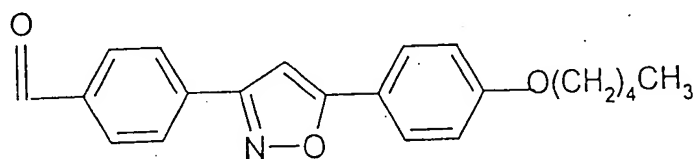
**Claim 5** (previously presented) A compound of claim 1 in which Y is hydrogen.

**Claim 6** (previously presented) A compound of claim 1 in which R<sub>3</sub> is methyl.

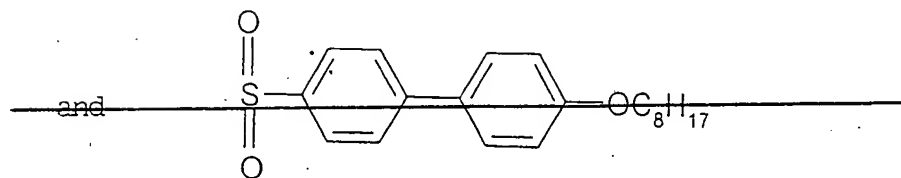
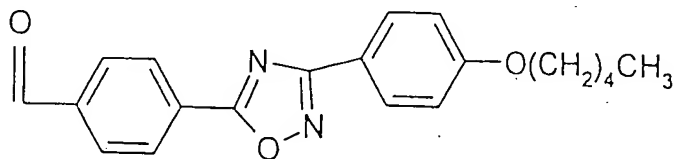
**Claim 7** (previously presented) A compound of claim 1 in which R<sub>4</sub> is hydroxyl.

**Claim 8** (currently amended) A compound of claim 1 in which R is selected from the group

consisting of

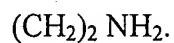


and

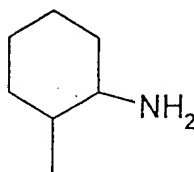


**Claim 9** (previously presented) A compound of claim 1 in which R<sub>1</sub> is hydrogen.

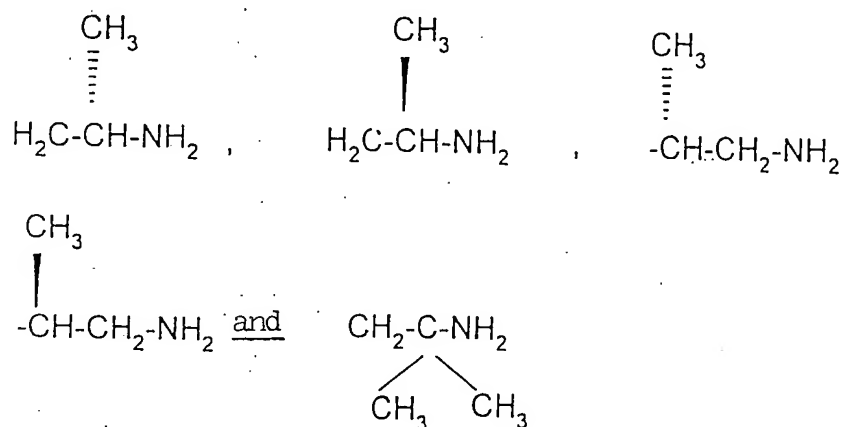
**Claim 10** (previously presented) A compound of claim 1 in which R<sub>2</sub> is



**Claim 11** (previously presented) A compound of claim 1 in which R<sub>2</sub> is



**Claim 12** (previously presented) A compound of claim 1 in which R<sub>2</sub> is selected from the group consisting of

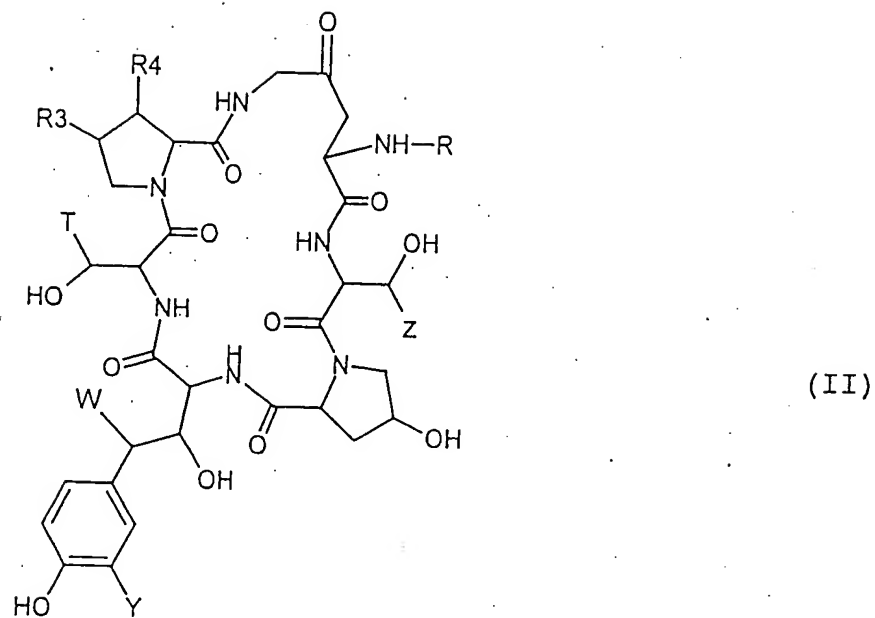


**Claim 13** (previously presented) A compound of claim 1 selected from the group consisting of

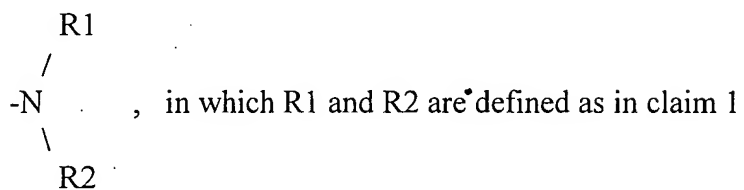
- 1-[4-[(2-aminoethyl)-amino]-N2-[[4-[5-[4-(pentyloxy)-phenyl]-3-isoxazolyl]-phenyl]-carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]-5-L-serine-echinocandine B trifluoroacetate,
- trans-1-[4-[(2-aminocyclohexyl)-amino]-N2-[[4-[5-[4-(pentyloxy)-phenyl]-3-isoxazolyl]-phenyl]-carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]-5-L-serine-echinocandine B trifluoroacetate,
- 1-[4-[(2(S)-aminopropyl)-amino]-N2-[[4-[5-[4-(pentyloxy)-phenyl]-3-isoxazolyl]-phenyl]-carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]-5-L-serine-echinocandine B trifluoroacetate,
- 1-[4-[(2-aminoethyl)amino]-N2-[[4-[5-[4-(pentyloxy)-phenyl]-1,3,4-thiadiazol-2-yl]-phenyl]-carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]-5-L-serine-echinocandine B trifluoroacetate,
- trans 1-[4-[(2-aminocyclohexyl)-amino]-N2-[[4-[5-[4-(pentyloxy)-phenyl]-1,3,4-thiadiazol-2-yl]-phenyl]-carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]-5-L-serine-echinocandine B trifluoroacetate and
- trans 1-[4-[(2-aminocyclohexyl)-amino]-N2-[[4-[3-[4-(pentyloxy)-phenyl]-1,2,4-oxadiazol-5-yl]-phenyl]-carbonyl]-L-ornithine]-4-[4-(4-hydroxyphenyl)-L-threonine]-5-L-serine-echinocandine B trifluoroacetate.

**Claim 14** (previously presented) A process for the preparation of a compound of claim 1

reacting a compound of the formula



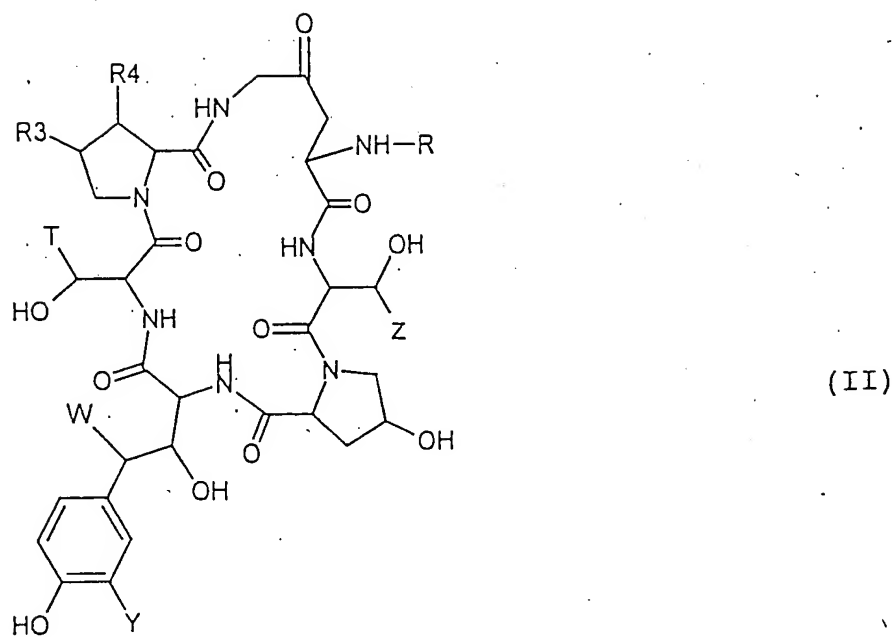
in which R, R<sub>3</sub>, R<sub>4</sub>, T, Y, W and Z are defined as in claim 1 with an amine or of an amine derivative capable of introducing



and optionally then with a reducing agent,  
and/or a functionalization agent of the amine,  
and/or an acid to form the salt of the product of claim 1,  
and/or a separation agent of the different isomers obtained.

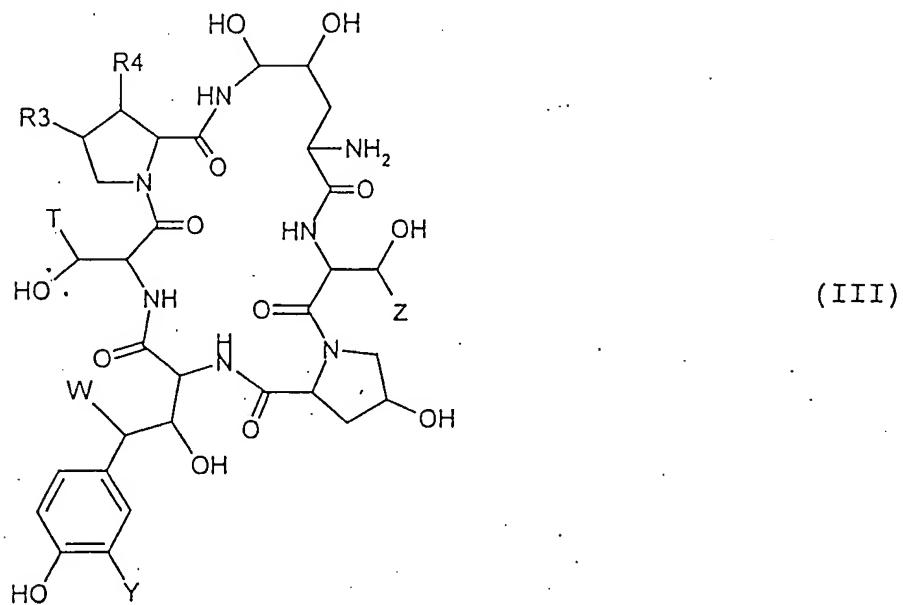
**Claim 15** (previously presented) A compound of the formula



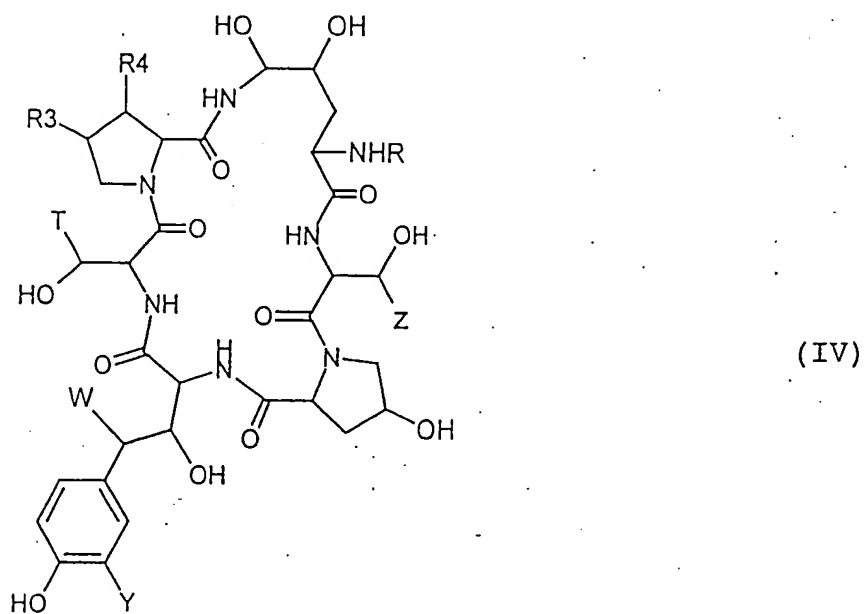


wherein R, R<sub>3</sub>, R<sub>4</sub>, T, Y, W and Z are defined as in claim 1.

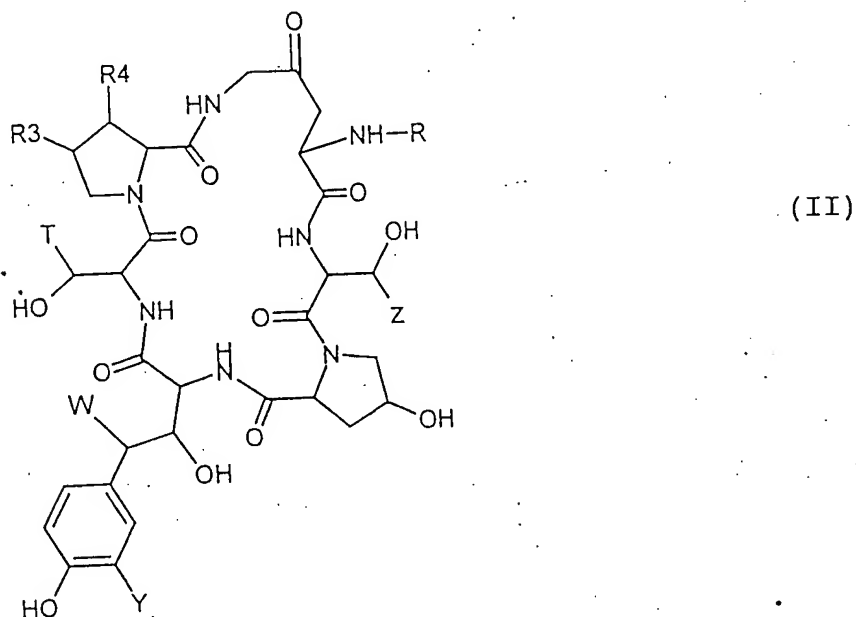
**Claim 16** (previously presented) A process of claim 14 wherein a compound of the formula



R<sub>3</sub>, R<sub>4</sub>, T, W, Y and Z are defined as in claim 14 reacted with an agent capable of replacing -NH<sub>2</sub> by -NHR, R being defined as in claim 14 to obtain a compound of the formula

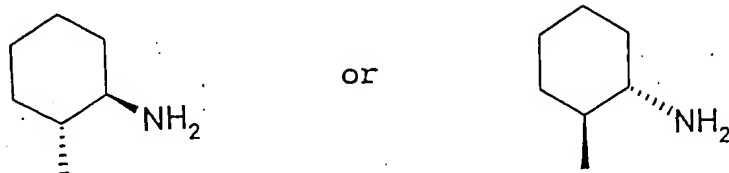


reacting the said compound with trimethylsilyl iodide to obtain the corresponding compound of the formula

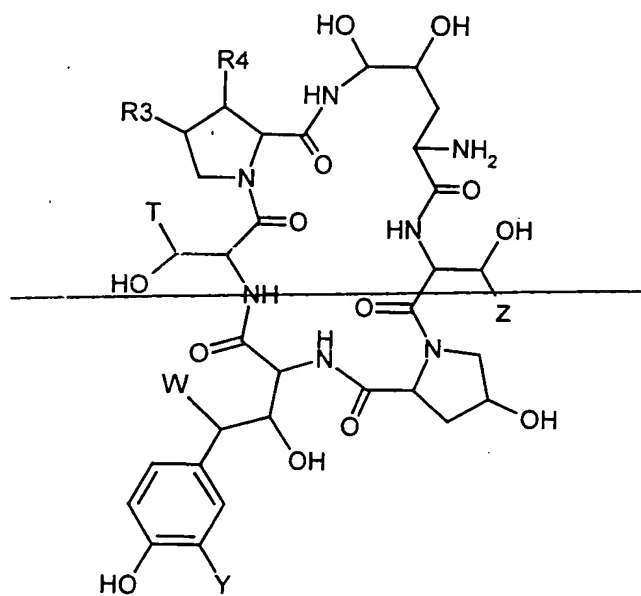


**Claims 17-19 (cancelled)**

**Claim 20** (previously presented) A compound of claim 11 wherein R<sub>2</sub> is

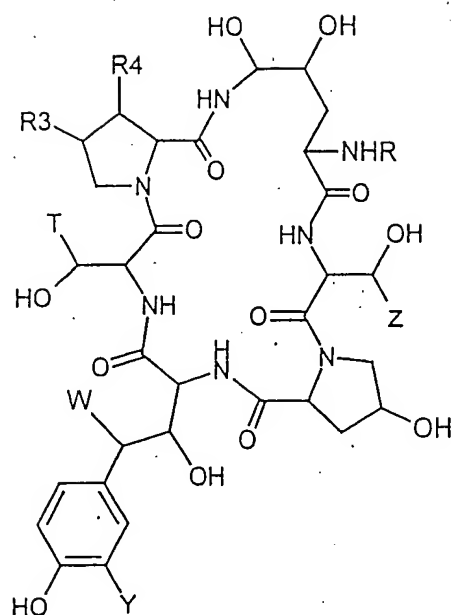


**Claim 21** (previously presented) A compound of the formula



(III)

~~or~~



(IV)

wherein R, R<sub>3</sub>, R<sub>4</sub>, T, W, Y and Z are defined as in claim 1.

**Claim 22** (currently amended) An antifungal composition comprising an antifungally effective amount of a compound of claim 1 and ~~in~~ an inert pharmaceutical carrier.

**Claim 23** (previously presented) A method of treating fungal infections in warm-blooded animals comprising administering to warm-blooded animals in need thereof an antifungally effective amount of a compound of claim 1.